

FILE 'HCAPLUS' ENTERED AT 08:22:52 ON 29 MAY 2007

L1 311474 S ANTIBODY
L2 4617 S (BETA-GLUCAN)
L3 754497 S CANCER OR TUMOR OR NEOPLAS?
L4 146686 S MONOCLONAL
L5 68113 S GD2 OR CD20 OR EGFR OR HER2 OR NEUROBLASTOMA OR MELANOMA OR L
L6 180 S L1 AND L2
L7 48 S L1 AND L2 AND L3
L8 22 S L1 AND L2 AND L3 AND L4
L9 7 S L1 AND L2 AND L3 AND L4 AND L5

FILE 'STNGUIDE' ENTERED AT 08:23:03 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:24:38 ON 29 MAY 2007

L10 81 S L6 AND (PY<2002 OR PRY<2002 OR AY<2002)
L11 18 S L7 AND (PY<2002 OR PRY<2002 OR AY<2002)
L12 3 S L8 AND (PY<2002 OR PRY<2002 OR AY<2002)
L13 1 S L9 AND (PY<2002 OR PRY<2002 OR AY<2002)

=> file hcaplus
COST IN U.S. DOLLARS
FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
0.84	0.84

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FILE COVERS 1907 - 29 May 2007 VOL 146 ISS 23
FILE LAST UPDATED: 28 May 2007 (20070528/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s antibody

L1 311474 ANTIBODY

=> s (beta-glucan)

1455756 BETA
14986 GLUCAN
L2 4617 (BETA-GLUCAN)
(BETA(W) GLUCAN)

=> s cancer or tumor or neoplas?

316729 CANCER
408979 TUMOR
494653 NEOPLAS?
L3 754497 CANCER OR TUMOR OR NEOPLAS?

=> s monoclonal

L4 146686 MONOCLONAL

=> s GD2 or CD20 or EGFR or HER2 or neuroblastoma or melanoma or lymphona or epidermoid

2208 GD2
3069 CD20
8590 EGFR
3603 HER2
16553 NEUROBLASTOMA
34665 MELANOMA
5 LYMPHONA
2337 EPIDERMOID
L5 68113 GD2 OR CD20 OR EGFR OR HER2 OR NEUROBLASTOMA OR MELANOMA OR
LYMPHONA OR EPIDERMOID

=> s L1 and L2

L6 180 L1 AND L2

=> s L1 and L2 and L3

L7 48 L1 AND L2 AND L3

=> s L1 and L2 and L3 and L4

L8 22 L1 AND L2 AND L3 AND L4

=> s L1 and L2 and L3 and L4 and L5

L9 7 L1 AND L2 AND L3 AND L4 AND L5

=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	2.60	3.44

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AND TECHNOLOGY CORPORATION, AND FACHINFORMATIONSZENTRUM KARLSRUHE

FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: May 25, 2007 (20070525/UP).

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.18	3.62

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FILE COVERS 1907 - 29 May 2007 VOL 146 ISS 23
FILE LAST UPDATED: 28 May 2007 (20070528/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s L6 and (PY<2002 or PRY<2002 or AY<2002)

21897330 PY<2002
3645664 PRY<2002

4174339 AY<2002
L10 81 L6 AND (PY<2002 OR PRY<2002 OR AY<2002)

=> s L7 and (PY<2002 or PRY<2002 or AY<2002)

21897330 PY<2002
3645664 PRY<2002
4174339 AY<2002
L11 18 L7 AND (PY<2002 OR PRY<2002 OR AY<2002)

=> s L8 and (PY<2002 or PRY<2002 or AY<2002)

21897330 PY<2002
3645664 PRY<2002
4174339 AY<2002
L12 3 L8 AND (PY<2002 OR PRY<2002 OR AY<2002)

=> s L9 and (PY<2002 or PRY<2002 or AY<2002)

21897330 PY<2002
3645664 PRY<2002
4174339 AY<2002
L13 1 L9 AND (PY<2002 OR PRY<2002 OR AY<2002)

=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	2.60	6.22

FILE 'STNGUIDE' ENTERED AT 08:24:51 ON 29 MAY 2007
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FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: May 25, 2007 (20070525/UP).

<-----User Break----->

=> d L13 ti abs bib
YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L13 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Antitumor antibody-enhancing glucan
AB This invention provides a composition comprising an effective amount of glucan capable of enhancing efficacy of antibodies. This invention further provides the above compns. and a pharmaceutically acceptable carrier. This invention also provides a method for treating a subject with cancer comprising administering the above-described composition comprising effective amount of glucan capable of enhancing efficacy of vaccines. This invention provides a composition comprising effective amount of glucan capable of enhancing efficacy of vaccines. This invention also provides a method of treating a subject comprising administering the above pharmaceutical composition to the subject. This invention provides a composition comprising effective amount of glucan capable of enhancing efficacy of natural antibodies. This invention provides a composition comprising effective amount of glucan capable of enhancing host immunity. This invention also provides a composition comprising effective amount of glucan capable of enhancing the action of an agent in preventing tissue rejection. It was shown that β -glucans greatly enhanced the antitumor effects of monoclonal antibodies against established

tumors in mice.
AN 2002:574940 HCAPLUS <<LOGINID::20070529>>
DN 137:119657
TI Antitumor antibody-enhancing glucan
IN Cheung, Nai-Kong V.
PA Sloan-Kettering Institute for Cancer Research, USA
SO PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	WO 2002058711	A1	20020801	WO 2002-US1276	20020115 <--	
	W:			AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW		
	RW:			GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		
	CA 2434938	A1	20020801	CA 2002-2434938	20020115 <--	
	AU 2002241905	A1	20020806	AU 2002-241905	20020115 <--	
	EP 1357919	A1	20031105	EP 2002-707502	20020115 <--	
	R:			AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR		
	US 2004116379	A1	20040617	US 2003-621027	20030716 <--	
	US 2006020128	A1	20060126	US 2005-218044	20050831 <--	
	US 2006160766	A1	20060720	US 2006-334763	20060117 <--	
PRAI	US 2001-261911P	P	20010116	<--		
	WO 2002-US1276	W	20020115			
	US 2003-621027	A1	20030716			
	WO 2004-US23099	A2	20040716			
	US 2005-218044	A2	20050831			

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d L12 1-3 ti
YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L12 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Antitumor antibody-enhancing glucan

L12 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Clostridial neurotoxin targeted conjugates for inhibition of secretion from non-neuronal cells

L12 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Polymeric drugs based on conjugates of synthetic and natural macromolecules. II. Anti-cancer activity of antibody or (Fab')₂-targeted conjugates and combined therapy with immunomodulators

=> d L12 2 3 ti abs bib
YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L12 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Clostridial neurotoxin targeted conjugates for inhibition of secretion from non-neuronal cells

AB A method of treatment of disease by inhibition of cellular secretory processes is provided. The method has particular application in the treatment of diseases dependent on the exocytotic activity of endocrine cells, exocrine cells, inflammatory cells, cells of the immune system, cells of the cardiovascular system, and bone cells. Agents and compns. therefor, as well as methods for manufacturing these agents and compns., are provided. In a preferred embodiment a clostridial neurotoxin, substantially devoid of holotoxin binding affinity for neuronal cells of the presynaptic muscular junction, is associated with a targeting moiety. The targeting moiety is selected such that the clostridial toxin conjugate so formed may be directed to a non-neuronal target cell to which the conjugate may bind. Following binding, a neurotoxin component of the conjugate, which is capable of inhibition of cellular secretion, passes into the cytosol of the target cell by cellular internalization mechanisms. Thereafter, inhibition of secretion from the target cell is effected.

AN 2001:228744 HCAPLUS <<LOGINID::20070529>>

DN 134:247267

TI Clostridial neurotoxin targeted conjugates for inhibition of secretion from non-neuronal cells

IN Foster, Keith Alan; Chaddock, John Andrew; Purkiss, John Robert; Quinn, Conrad Padraig

PA Microbiological Research Authority, UK

SO PCT Int. Appl., 63 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001021213	A2	20010329	WO 2000-GB3669	20000925 <--
	WO 2001021213	A3	20020711		
	W:				
	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
	RW:				
	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2383470	A1	20010329	CA 2000-2383470	20000925 <--
	EP 1235594	A2	20020904	EP 2000-962721	20000925 <--
	R:				
	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				
	JP 2003509476	T	20030311	JP 2001-524636	20000925 <--
	AU 782457	B2	20050728	AU 2000-74365	20000925 <--
	US 2003180289	A1	20030925	US 2002-88665	20020814 <--
	AU 2005227383	A1	20051124	AU 2005-227383	20051027 <--
	US 2006216283	A1	20060928	US 2006-327855	20060109 <--
PRAI	GB 1999-22554	A	19990923	<--	
	WO 2000-GB3669	W	20000925	<--	
	WO 2000-GB3681	A	20000925	<--	
	US 2002-88665	A1	20020814		

L12 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Polymeric drugs based on conjugates of synthetic and natural macromolecules. II. Anti-cancer activity of antibody or (Fab')₂-targeted conjugates and combined therapy with immunomodulators

AB We provide data on in vivo targeting of the Thy 1.2 (CDw90) cell surface receptor expressed on neoplastic T cells, mouse EL4 T cell lymphoma. The targeting antibody and the anticancer drug,

doxorubicin (DOX) were conjugated to a water-soluble copolymer based on N-(2-hydroxypropyl)methacrylamide (HPMA) acting as a carrier responsible for controlled intracellular release of the conjugated drug. The in vivo therapeutic efficacy of HPMA copolymer-bound DOX targeted with anti-EL4 antibody, polyclonal anti-thymocyte globulin (ATG), monoclonal anti-Thy 1.2 antibody or its F(ab')₂ fragment was compared with the efficacy of DOX conjugated to HPMA copolymer containing nonspecific IgG or bovine serum albumin (BSA). Anti-EL4 antibody-targeted conjugate caused a significant retardation of tumor growth and an extension of the life span of treated mice. The effect was comparable with that of HPMA copolymer-bound DOX targeted with ATG, anti-Thy 1.2 antibody or its F(ab')₂ fragment. However, considerable antitumor effect was seen also in conjugates targeted instead of specific antibodies with syngeneic nonspecific IgG or BSA. Patients with advanced cancer are often immunocompromised due to dysfunction of their immune system induced by cancer and cytotoxic drugs. A significant decrease of unwanted side-effects of targeted drugs against a number of vital organs was already documented. In this study we have compared immunotoxic effects of free DOX with those of its antibody-targeted form on NK cells and cytolytic T lymphocytes (CTLs) isolated from C57BL/10 mice bearing EL4 T cell lymphoma. In the same model we have tested the combination therapy with immunomodulators (β-glucan or AM-2) injected together with targeted daunomycin. We have observed a significant protective effect of targeted DOX against NK cells and CTLs. Moreover, the data revealed that combination therapy considerably enhances antitumor efficacy of the targeted anticancer drug.

AN 2000:46595 HCAPLUS <<LOGINID::20070529>>
 DN 132:284054
 TI Polymeric drugs based on conjugates of synthetic and natural macromolecules. II. Anti-cancer activity of antibody or (Fab')₂-targeted conjugates and combined therapy with immunomodulators
 AU Rihova, B.; Jelinkova, M.; Strohal, J.; Subr, V.; Plocova, D.; Hovorka, O.; Novak, M.; Plundrova, D.; Germano, Y.; Ulbrich, K.
 CS Institute of Microbiology, Academy of Sciences of the Czech Republic, Prague, 142 20, Czech Rep.
 SO Journal of Controlled Release (2000), 64(1-3), 241-261
 CODEN: JCREEC; ISSN: 0168-3659
 PB Elsevier Science Ireland Ltd.
 DT Journal
 LA English
 RE.CNT 49 THERE ARE 49 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d l11 1-18 ti

YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L11 ANSWER 1 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Therapy-enhancing glucan

L11 ANSWER 2 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Macrophage receptor Dectin-1

L11 ANSWER 3 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Antitumor antibody-enhancing glucan

L11 ANSWER 4 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Plants, polysaccharides, and the treatment and prevention of neoplasia

L11 ANSWER 5 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Clostridial neurotoxin targeted conjugates for inhibition of secretion from non-neuronal cells

L11 ANSWER 6 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Preparation of antibody against antitumor β - glucan in Grifola frondosa and its application

L11 ANSWER 7 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Immunopharmacological and immunotoxicological activities of a water-soluble (1 \rightarrow 3)- β -D-glucan, CSBG from Candida spp

L11 ANSWER 8 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Failure in antitumor activity by overdose of an immunomodulating . beta.-glucan preparation, sonifilan

L11 ANSWER 9 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Polymeric drugs based on conjugates of synthetic and natural macromolecules. II. Anti-cancer activity of antibody or (Fab')₂-targeted conjugates and combined therapy with immunomodulators

L11 ANSWER 10 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Antigen-specific response of murine immune system toward a yeast . beta.-glucan preparation, zymosan

L11 ANSWER 11 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Activation of murine macrophages by grifolan

L11 ANSWER 12 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Cellular requirements for immunomodulatory effects caused by cell wall components of Paracoccidioides brasiliensis on antibody production

L11 ANSWER 13 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Preparation and specificity of antibodies to an anti-tumor . beta.-glucan, lentinan

L11 ANSWER 14 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Covalently bound β -glucan conjugates with bioactive agents for targeted delivery

L11 ANSWER 15 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Interrelation of structure and antitumor effects of fungal (1 \rightarrow 3) β -D-glucans.

L11 ANSWER 16 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Pulmonary metastases neutralization and tumor rejection by in vivo administration of β glucan and bispecific antibody

L11 ANSWER 17 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Straw mushroom, fukurotake, Volvariella volvacea

L11 ANSWER 18 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Antitumor and immunomodulating activities of a β - glucan obtained from liquid-cultured Grifola frondosa

=> d l11 4 6 7 8 10 11 13 14 16 17 18 ti abs bib

YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L11 ANSWER 4 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Plants, polysaccharides, and the treatment and prevention of

neoplasia

AB A review. Plants and Fungi have traditionally been the single largest source of lead compds. for the development of therapeutics by the pharmaceutical industry. Currently mushroom and plant polysaccharides brought to attention by Complementary and Alternative medicine, are undergoing scientific anal. and development to prevent and treat cancer. Two classes of saccharides are under investigation- beta glucan polysaccharides as biol. response modifiers for the adjuvant treatment of cancer and "Oligosaccharin"- related oligosaccharides for the prevention of sun-induced skin cancer. Beta glucans already in human trials in the Far East will require mechanistic pharmacol. studies and definition of structure function relationships before they are ready for clin. trials in the West. Other beta glucans that prime natural killer cells for antibody dependent cell-mediated cytotoxicity are approaching clin. trials. Oligosaccharides that downregulate production of immuno-suppressive cytokines by UV radiation injured keratinocytes are promising agents for the prevention of environmental skin cancer.

AN 2001:398732 HCAPLUS <<LOGINID::20070529>>

DN 136:160666

TI Plants, polysaccharides, and the treatment and prevention of neoplasia

AU Pelley, Ronald P.; Strickland, Faith M.

CS Pangea Phytoceuticals, Harlingen, TX, 78550, USA

SO Critical Reviews in Oncogenesis (2000), 11(3&4), 189-225

CODEN: CRONEI; ISSN: 0893-9675

PB Begell House, Inc.

DT Journal; General Review

LA English

RE.CNT 197 THERE ARE 197 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 6 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Preparation of antibody against antitumor β - glucan in Grifola frondosa and its application

AB Antibodies against an antitumor β -glucan purified from Grifola frondosa (GGF) were raised in the rabbit by s.c. immunization. Our antibodies reacted significantly with GGF by an ELISA inhibition assay. The antibodies did not recognize other polysaccharides such as laminarin and pustulan, but reacted somewhat with lentinan, whose structure is similar to GGF. It was demonstrated that GGF could be measured by ELISA using antibodies. In addition, the effects of the storage temperature on GGF content during storage were measured using our antibody. GGF content was 24.7 μ g/g fresh weight (f.w.) at zero time storage, and little change occurred during storage of the mushroom for 7 days at 5°. However, a drastic decrease to 11.4 μ g/g f.w. occurred after 7 days of storage at 20°. These results suggest that storage at low temps. is desirable to maintain the quality of GGF.

AN 2000:308382 HCAPLUS <<LOGINID::20070529>>

DN 133:320973

TI Preparation of antibody against antitumor β - glucan in Grifola frondosa and its application

AU Mizuno, Masashi; Yamakawa, Akio; Minato, Ken-Ichiro; Kawakami, Sachiko; Tatsuoka, Shigenobu; Terai, Hirofumi; Tsuchida, Hironobu

CS Graduate School of Science and Technology, Kobe University, Kobe, 657-8501, Japan

SO Food Science and Technology Research (1999), 5(4), 398-401

CODEN: FSTRFS; ISSN: 1344-6606

PB Japanese Society for Food Science and Technology

DT Journal

LA English

RE.CNT 22 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 7 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Immunopharmacological and immunotoxicological activities of a
 water-soluble (1 → 3)-β-D-glucan, CSBG from *Candida* spp
 AB We have established a convenient, two-step procedure to solubilize the
 yeast cell wall (1→3)-β-D-glucan using the combination of
 NaClO oxidation and DMSO extraction. *Candida* soluble β-D-glucan (CSBG) was
 mainly composed of a linear β-1,3 glucan with a linear
 β-1,6-glucan moiety. In this study, we screened for several
 immunopharmacol. activities of CSBG and found the following activities:
 (1) interleukin-6 synthesis of macrophages in vitro; (2) antagonistic
 effect for zymosan mediated-tumor necrosis factor synthesis of
 macrophages; (3) augmentation for lipopolysaccharide mediated
 tumor necrosis factor and nitrogen oxide syntheses of macrophages;
 (4) activation of alternative pathway of complement; (5) hematopoietic
 response on cyclophosphamide induced leukopenia; (6) the antitumor effect
 on ascites form tumor; (7) Enhanced vascular permeability; (8)
 priming effect on lipopolysaccharide triggered TNF-α synthesis; and
 (9) adjuvant effect on antibody production. These results strongly
 suggested that CSBG possessed various immunopharmacol. activity.
 AN 2000:235041 HCAPLUS <<LOGINID::20070529>>
 DN 133:12504
 TI Immunopharmacological and immunotoxicological activities of a
 water-soluble (1 → 3)-β-D-glucan, CSBG from *Candida* spp
 AU Tokunaka, Kazuhiro; Ohno, Naohito; Adachi, Yoshiyuki; Tanaka, Shigenori;
 Tamura, Hiroshi; Yadomae, Toshiro
 CS Laboratory for Immunopharmacology of Microbial Products, School of
 Pharmacy, Tokyo University of Pharmacy and Life Science, Tokyo, 192-0392,
 Japan
 SO International Journal of Immunopharmacology (2000), 22(5),
 383-394
 CODEN: IJIMDS; ISSN: 0192-0561
 PB Elsevier Science Ltd.
 DT Journal
 LA English
 RE.CNT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
 TI Failure in antitumor activity by overdose of an immunomodulating .
 beta.-glucan preparation, sonifilan
 AB Schizophyllan (SPG, Sonifilan) is a soluble (1→3)-β-D-glucan,
 used as a biol. response modifier (BRM) with radiation therapy for
 cancer treatment in Japan. The mechanism of SPG-mediated
 antitumor activity is thought to be via immune stimulation, which includes
 cytokine production, hematopoietic response, and so on. In this paper, we
 found that the activity of SPG was quite long-lived and an overdose
 significantly failed to display the antitumor activity. To demonstrate
 the mechanism several parameters were examined using a high dose of SPG
 administration as follows: i) the effect on vascular permeability in vivo,
 ii) the priming effect on tumor necrosis factor (TNF-α)
 production in vivo, iii) the effect on macrophage adherence to plastic plate
 in vitro, and iv) anti-Sarcoma 180 antibody production in vivo. It
 was evident that vascular permeability and anti-Sarcoma 180
 antibody production remained unchanged, but TNF-α production and
 adherence to a plastic plate was significantly reduced by a high dose of
 SPG. These facts strongly suggested that modulation of the cytokine
 syntheses and the leukocyte traffic would be the causative mechanisms of
 the failure of antitumor activity by an overdose of SPG.
 AN 2000:97854 HCAPLUS <<LOGINID::20070529>>
 DN 132:245973
 TI Failure in antitumor activity by overdose of an immunomodulating .
 beta.-glucan preparation, sonifilan
 AU Miura, Toshihide; Miura, Noriko N.; Ohno, Naohito; Adachi, Yoshiyuki;
 Shimada, Shigehiko; Yadomae, Toshiro

CS Laboratory for Immunopharmacology of Microbial Products, School of
Pharmacy, Tokyo University of Pharmacy and Life Science, Tokyo, 192-0392,
Japan

SO Biological & Pharmaceutical Bulletin (2000), 23(2), 249-253
CODEN: BPBLEO; ISSN: 0918-6158

PB Pharmaceutical Society of Japan

DT Journal

LA English

RE.CNT 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 10 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Antigen-specific response of murine immune system toward a yeast .
beta.-glucan preparation, zymosan

AB Zymosan, a particulate β -glucan preparation from
Saccharomyces cerevisiae, shows various biol. activities, including anti-
tumor activity. We have previously shown that soluble .beta
.-glucan initiated anti-tumor activity was long-lived
and was effective even by prophylactic treatment at 1 mo prior to
tumor challenge. However, the activity by zymosan was relatively
short-lived. Antigen-specific responses of mice to zymosan might be a
causative mechanism. In this paper, mice were immunized with zymosan and
antibody production and antigen-specific responses of lymphocytes to
zymosan were analyzed. Sera of zymosan immune mice contained
zymosan-specific IgG assessed by ELISA and FACS. Spleen and bone marrow
cells of zymosan-immune mice showed higher cytokine production in response to
zymosan. Specificity of zymosan-specific responses were also analyzed
using various derivs. prepared from zymosan. These facts strongly suggested
that mice recognize zymosan as antigen in addition to non-specific immune
stimulant.

AN 1999:311543 HCAPLUS <<LOGINID::20070529>>

DN 131:128740

TI Antigen-specific response of murine immune system toward a yeast .
beta.-glucan preparation, zymosan

AU Miura, T.; Ohno, N.; Miura, N. N.; Adachi, Y.; Shimada, S.; Yadomae, T.

CS School of Pharmacy, Laboratory for Immunopharmacology of Microbial.
Products, Tokyo University of Pharmacy and Life Science, Hachioji, Tokyo,
192-0392, Japan

SO FEMS Immunology and Medical Microbiology (1999), 24(2), 131-139
CODEN: FIMIEV; ISSN: 0928-8244

PB Elsevier Science B.V.

DT Journal

LA English

RE.CNT 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 11 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Activation of murine macrophages by grifolan

AB A gel-forming (1 \rightarrow 3)- β -D-glucan, grifolan (GRN) from an edible
mushroom (*Grifola frondosa*), enhances various immunol. activities. Here,
effect of GRN on the induction of cytokines and nitric oxide by macrophage
(MP) cell line (RAW264.7), peritoneal MP (PM), and Kupffer cell is shown.
GRN bound to MP was detected immunohistochem., using an anti-GRN
antibody. GRN could induce production of TNF α , IL-1 α , and
IL-6 by RAW264.7. Incubation with GRN also induced those cytokines in PM.
GRN induced phosphorylation of MAP kinase and p38 of PM. The kinetic
study on the activation of Kupffer cells revealed that GRN could induce
enhanced production of cytokines and nitric oxide on days 4-7 after i.v.
administration of GRN. Cytostatic activity of Kupffer cells against
murine lymphoma, EL-4, was also augmented by GRN with similar time course
to nitric oxide production. The cytostatic activity was dependent on nitric
oxide, since an iNOS inhibitor diminished the cytostatic activity.
Administration of GRN increased expression of CD11b, known as the .
beta.-glucan receptor, on Kupffer cells on day 7.

Apparently, GRN can activate murine MPs to enhance production of cytokines and nitric oxide.

AN 1998:453248 HCAPLUS <<LOGINID::20070529>>

DN 129:211409

TI Activation of murine macrophages by grifolan

AU Adachi, Y.; Takano, E.; Ohno, N.; Yadomae, T.

CS School of Pharmacy, Tokyo University of Pharmacy and Life Science, Tokyo, Japan

SO Proceedings - Beltwide Cotton Conferences (1998), (Vol. 1), 262-266

CODEN: PCOCEN; ISSN: 1059-2644

PB National Cotton Council

DT Journal

LA English

RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 13 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Preparation and specificity of antibodies to an anti-tumor .
beta.-glucan, lentinan

AB Antibodies against β -glucan, lentinan from "Shiitake" (*Lentinus edodes*), were raised in the rabbit by s.c. immunization. Our antibodies reacted significantly with lentinan by inhibition assay of ELISA. The antibodies did not recognize the other polysaccharides such as amylose, dextran, laminarin and galactan. It was proved that lentinan contents in mushroom could be measured by ELISA with the anti-lentinan antisera. Its contents were 3.5 mg/g fresh weight in *Lentinus edodes*. However, lentinan was not contained in *Agaricus brazei*, *Agaricus bisporus* and *Romania bitrytis*.

AN 1997:90871 HCAPLUS <<LOGINID::20070529>>

DN 126:170161

TI Preparation and specificity of antibodies to an anti-tumor .
beta.-glucan, lentinan

AU Mizono, Masashi; Minato, Ken-ichiro; Tsuchida, Hironobu

CS Grad. Sch. Sci. and Tech., Kobe Univ., Kobe, 657, Japan

SO Biochemistry and Molecular Biology International (1996), 39(4), 679-685

CODEN: BMBIES; ISSN: 1039-9712

PB Academic

DT Journal

LA English

L11 ANSWER 14 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Covalently bound β -glucan conjugates with
bioactive agents for targeted delivery

AB A glucan composition is disclosed which contains a β -1,3-glucan covalently attached to a bioactive agent. The β -1,3-glucan is attached to the bioactive agent by means of a hydrolyzable covalent linkage to form a glucan/agent complex. Also disclosed are methods relating to the complex of the invention, including a method for the treatment of a pathogen capable of invading or colonizing phagocytic cells, and a method for delivering an antigen to a phagocytic cell. Purification of glucan from *Euglena gracilis* is described. Also described is e.g. preparation of a β -1,3-glucan conjugate with herpes simplex virus gD2 glucoprotein. The conjugate had enhanced adjuvant activity.

AN 1996:462438 HCAPLUS <<LOGINID::20070529>>

DN 125:105156

TI Covalently bound β -glucan conjugates with
bioactive agents for targeted delivery

IN Tuse, Daniel; Mohagheghpour, Nahid; Dawson, Marcia; Hobbs, Peter; Winant, Richard

PA Sri International, USA

SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2

DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9614873	A2	19960523	WO 1995-US14800	19951114 <--
	WO 9614873	A3	19960829		
	W: CA, JP				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
PRAI	US 1994-340831	A	19941116 <--		

L11 ANSWER 16 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Pulmonary metastases neutralization and tumor rejection by in vivo administration of β glucan and bispecific antibody

AB Bispecific antibody (BsAb) with specificity for tumor cell surface antigen and the CD3 mol. on T cells can redirect activated T cells to lyse tumor cells. Since the ex vivo expansion and activation of T cells is impractical and ineffective for treating established tumors, the authors tested whether the immune stimulant . beta. glucan could in situ-activate T cells, which could secondarily be retargeted with BsAbs to lyse tumor cells. To test for tumor neutralization, C3H/HeN mice were injected i.v. with C1-62 melanoma cells and immediately treated with i.p. .beta. glucan and/or anti-CD3 (500A2) + anti-p97 (96.5) F(ab')₂ BsAb i.v. Pulmonary metastases were counted 14 days later. To test for tumor rejection and survival in a solid tumor model, mice were injected s.c. and i.p. with C1-62 cells and 7 days later administered β glucan i.p. and/or F(ab')₂ BsAb i.v. In the neutralization model, there was a significant reduction in the number of metastases in the β glucan + BsAb group, as compared with controls, and with β glucan alone. In the established tumor model, β glucan + BsAb reduced the incidence of s.c. tumors as compared with control, BsAb alone, and β glucan alone. It also prolonged survival of tumor-bearing mice compared with control, BsAb alone, and β glucan alone. Thus, T cells can be activated in vivo by β glucan and retargeted with F(ab')₂ BsAb.

AN 1996:160223 HCAPLUS <<LOGINID::20070529>>

DN 124:257967

TI Pulmonary metastases neutralization and tumor rejection by in vivo administration of β glucan and bispecific antibody

AU Penna, Christophe; Dean, Phillip A.; Nelson, Heidi

CS Department Surgery, Mayo Clinic and Mayo Foundation, Rochester, MN, 55905, USA

SO International Journal of Cancer (1996), 65(3), 377-82
CODEN: IJCNAW; ISSN: 0020-7136

PB Wiley-Liss

DT Journal

LA English

L11 ANSWER 17 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN

TI Straw mushroom, fukurotake, Volvariella volvacea

AB A review with 14 listed refs. on the systematic fractionation and structural diversity of branched (1 \rightarrow 3)- β -glucan of fukurotake, chemical modification in relation to immunomodulating mechanism of the glucans, antibodies to the glucans and their application in studies of neoplasm inhibition.

AN 1995:536205 HCAPLUS <<LOGINID::20070529>>

DN 123:141915

TI Straw mushroom, fukurotake, Volvariella volvacea

AU Misaki, Akira; Kishida, Etsu

CS Osaka City University, Ashiya, 659, Japan
SO Food Reviews International (1995), 11(1), 219-23
CODEN: FRINEL; ISSN: 8755-9129
DT Journal; General Review
LA English

L11 ANSWER 18 OF 18 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Antitumor and immunomodulating activities of a β -
glucan obtained from liquid-cultured Grifola frondosa
AB The effects of the β -1,3-glucan, LELFD, obtained from liquid-cultured
mycelium of G. frondosa, on the growth of syngeneic tumors and immune
responses in mice were examined. In Meth A fibrosarcoma or IMC carcinoma
solid tumor systems, LELFD administered i.p. or intralesionally
(i.l.) exhibited significant antitumor effects. However, the growth of
L1210 and P388 leukemias was unaffected by the injection of LELFD. The
injection of LELFD i.p. enhanced the activities of natural killer cells
and macrophages in mice. LELFD also enhanced the antibody
response when it was injected i.p. with sheep red blood cells into mice.
Furthermore, it was found that LELFD could activate complement pathway.
AN 1989:185485 HCAPLUS <<LOGINID::20070529>>
DN 110:185485
TI Antitumor and immunomodulating activities of a β -
glucan obtained from liquid-cultured Grifola frondosa
AU Suzuki, Iwao; Hashimoto, Koichi; Oikawa, Shozo; Sato, Kichiro; Osawa,
Masumi; Yadomae, Toshiro
CS Tokyo Coll. Pharm., Hachioji, 192-03, Japan
SO Chemical & Pharmaceutical Bulletin (1989), 37(2), 410-13
CODEN: CPBTAL; ISSN: 0009-2363
DT Journal
LA English

=> file caplus

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	0.06	66.52
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
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FILE LAST UPDATED: 28 May 2007 (20070528/ED)

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=> s l11 and oral
311474 ANTIBODY
1455756 BETA
14986 GLUCAN
4617 (BETA-GLUCAN)
(BETA(W) GLUCAN)
316729 CANCER
408979 TUMOR
494653 NEOPLAS?
21897330 PY<2002
3645664 PRY<2002
4174339 AY<2002
207085 ORAL
L14 1 L11 AND ORAL

=> d l14 ti

L14 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN
TI Therapy-enhancing glucan

=> d his

(FILE 'HOME' ENTERED AT 08:20:33 ON 29 MAY 2007)

FILE 'HCAPLUS' ENTERED AT 08:22:52 ON 29 MAY 2007

L1 311474 S ANTIBODY
L2 4617 S (BETA-GLUCAN)
L3 754497 S CANCER OR TUMOR OR NEOPLAS?
L4 146686 S MONOCLONAL
L5 68113 S GD2 OR CD20 OR EGFR OR HER2 OR NEUROBLASTOMA OR MELANOMA OR L
L6 180 S L1 AND L2
L7 48 S L1 AND L2 AND L3
L8 22 S L1 AND L2 AND L3 AND L4
L9 7 S L1 AND L2 AND L3 AND L4 AND L5

FILE 'STNGUIDE' ENTERED AT 08:23:03 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:24:38 ON 29 MAY 2007

L10 81 S L6 AND (PY<2002 OR PRY<2002 OR AY<2002)
L11 18 S L7 AND (PY<2002 OR PRY<2002 OR AY<2002)
L12 3 S L8 AND (PY<2002 OR PRY<2002 OR AY<2002)
L13 1 S L9 AND (PY<2002 OR PRY<2002 OR AY<2002)

FILE 'STNGUIDE' ENTERED AT 08:24:51 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:25:06 ON 29 MAY 2007

FILE 'STNGUIDE' ENTERED AT 08:25:07 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:25:17 ON 29 MAY 2007

FILE 'STNGUIDE' ENTERED AT 08:25:17 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:25:34 ON 29 MAY 2007

FILE 'STNGUIDE' ENTERED AT 08:25:35 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:26:36 ON 29 MAY 2007

FILE 'STNGUIDE' ENTERED AT 08:26:37 ON 29 MAY 2007

FILE 'HCAPLUS' ENTERED AT 08:28:09 ON 29 MAY 2007

FILE 'STNGUIDE' ENTERED AT 08:28:10 ON 29 MAY 2007

FILE 'CAPLUS' ENTERED AT 08:28:15 ON 29 MAY 2007

L14 1 S L11 AND ORAL

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FULL ESTIMATED COST	20.91	87.43
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	ENTRY	SESSION
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	ENTRY	SESSION
FULL ESTIMATED COST	20.91	87.43
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-10.92

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	21.38	87.90
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-10.92

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=> s oral

L15 207085 ORAL

=> s L10 and L15

L16 3 L10 AND L15

=> file stnguide

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FULL ESTIMATED COST	2.60	90.50
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=> d l16 1-3 ti

YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L16 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Therapy-enhancing glucan

L16 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Immortal cell line derived from the grouper *Epinephelus coioides* and the applications thereof

L16 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Treatment of fungal infections with polyene or beta glucan synthase inhibitor antifungals combined with anti HSP90 antibodies

=> d l16 2 3 ti abs bib

YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L16 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN
TI Immortal cell line derived from the grouper *Epinephelus coioides* and the applications thereof
AB The invention comprises the generation of antibodies against nervous necrosis virus (NNV) and infectious pancreatic necrosis (IPNV) virus. The